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(54) COSMETIC COMPOSITIONS

(71) We, HENKEL & CIF, GMBH, a German Company, of 67, Henkeisurasse, Duesseldorf, 4000, Germany, do hereby declare the invention, for which we pray that a patent may be granted to us, and the method by which it is to be performed, to be particularly described in and by the following statement:—

The invention relates to a composition for use in cosmetic preparations, especially cosmetic cleansing agents, containing substances for replacing the oil in the skin, based on esterification products of glycerine-ethylene oxide adducts with long-chain fatty solds.

Cosmetic cleansing agents such as shampoos, form baths, toilet soaps and similar products cause a more or less great removal of oil from the skin on repeated use. This phenomenon is particularly pronounced when the cleansing agents are based on synthetic, surface-active substances such as alkylbenzene sulphonates, fatty alcohol sulphates, olefine sulphonates, fatty alcohol ether suiphates and other surfaceactive compounds. Attempts have therefore been made to neutralise this removal of oil from the skin by replacing the oil by means of suitable additions to the cleansing agents. Allowance has in such case to be made for disadvantages, however, since the products 30 generally concerned in replacing the oil in the skin have an unfavourable action on the foaming properties of the cleansing agents and, in cosmetic preparations based on alcoholwater mixtures, show insufficient solubility.

The present invention provides a composition for use in cosmetic preparation which comprises a surface active compound and the esterification product of an ethylene oxide addition compound produced from glycerine and 4 to 20 mul of ethylene oxide per mol of glycerine, with a fatty acid of chain length from 8 to 18 carbon atoms in a ratio of 1 to 2 mol of fatty acid to 1 mol of glycerine-ethylene oxide addition compound as oil replacement material.

Esterification products of ethylene oxide addition compounds produced from glycerine and 7 to 15 mol of ethylene oxide per mol of glycerine with fatty acids of chain length from 8 to 18 carbon atoms in a ratio of 1 mol of fatty acid to 1 mol of glycerine-ethylene oxide addition compound are preferred as the oil replacement materials.

The preparation of the ethylene oxide addition compound as intermediate product was generally effected in known way by reacting glycerine with ethylene oxide in the desired proportions with elkaline catalysis by means of sodium ethylate. For the further treatment, the ethylene oxide addition compound obtained was reacted in the usual way with a fatty acid of chain length from 8 to 18 carbon atoms in the molar ratio of 1:1 or 1:2, using isopropyl titunate as esterification catalyst. The esterification products obtained were light-coloured to yellowish liquids of low viscosity with an oil character to land-like products of a faint self colour.

The quantities of oil replacement material according to the invention used in the cosmetic preparations may vary within very wide limits according to the product and its oil-removing action, and generally vary from 2 to 50% by weight, especially 5 to 25% by weight. Still higher additions are possible if the esterification products according to the invention are used at the same time in their property as surface-active substances, but in most cases this use will be of small advantage.

Oil replacement materials to be used according to the invention include, for example, esterification products from

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÷	the adduct of	1 mol c	f glyceri	ne + 4 mol of citylene oxide with 1 mol of coconut fatty acid C _{8-a8}
	23.	19	>>	+6 mol of enhylene oxide with 2 mol of coconut fatty acid C ₈₋₁₈
5	59	32	72	47 mol of ethylene oxide with 1 mol of excount fatty acid C ₈₋₁₈
	32	28	59	+7 mol of ethylene oxide with 1 mol of rallow farty acid
10	59	\$7	35	+8 mol of ethylene oxide with 1 mol of oleic acid
	32	23	518	+ 9 mol of ethylene oxide with 1 mol of palm kernel fatry acid
	33	35	,>>	+ 10 mol of ethylene oxide with 1 mol of tallow fatty acid
15	3>	,25	53	+ 10 mol of ethylene oxide with 2 mol
	232	59	,55	4 12 mol of ethylene oxide with 1 mol of coconut fatty acid C ₈₋₁₈
23	**	39	23	+ 15 mol of ethylene oxide with 1 mol of palm kernel fatty acid
	r,c	22	25	+ 15 mol of ethylene oxide with 2 mol of tellow faity acid

The present invention will be further described by way of illustration with reference to the following examples. Abbreviations and units used in the examples are defined as follows: -

"Acid value" is the number of mg of potassium hydroxide which are needed to neutralize the free esterified fatty acid contained in I g of substance;

"Saponification value" is the number of mg of potassium hydroxide which are needed to

completely saponify 1 g of ester; "Hydroxyl number" is the number of mg of porassium hydroxide which are needed to neutralize the acetic acid which is absorbed by 1 g of material. The procedure used is as follows; the substance being investigated is weighed out exactly and is then acetylated by heating with crystalline acetic acid, all the OH groups present thereby being esterified. Subsequently, the acctylated portion of the material is separated by a shaking and washing pro-45 cedure and the acetylated part is saponified using boiling potassium hydroxide solution. The excess potassium hydroxide which is pre-

sent after the saponification steps is determined

by attration with hydrochloric acid.

50 "EO" is the ethylene oxide group; "WAS" is active washing substance, All parts used throughout the examples are by weight unless otherwise specified.

Examples The following esterification products were used for the experiments and cosmeric prepararions described below.

(1 mol of glycerine + 7.4 mol of ethylene oxide) with 1 mol of coconut fatty acid Acid value 1.0, sap. value 92, hydroxy value 185

(1 mol glycerine + 7.4 mol of ethylene oxide) with I mol of tallow fatty acid (Acid value 1.1, sap. value 83, hydroxy value 166

(1 mol glycerine + 10 mol of ethylene oxide) with 1 mol of tallow fatty acid Acid value 1.4, sap. value 71, hydroxy

Since for cosmetic cleansing compositions the ability to combine with certain surface-active compounds is of essential importance, mixtures given in the following Table were tested.

TABLE I

Mixing component	Mixture 1	Mixture 2	Mixture 3
(A)	10	***************************************	••••
(B)		10	••••
(C)	••••	Arres	10
Sodium lauryl ether sulphate (2 EO) (27—28% WAS)	50	50	50
Water	40	40	40
Result	Clear homo- geneous solution	Clear homo- geneous solution	Clear homo- geneous solution

In a further experiment the foaming power the individual oil replacement materials was of a foam bath basic recipe with additions of examined.

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TABLE II

Components	Mixture 1	Mixture 2	Mixture 3	Mixture 4	Mixture 5
Sodium Isuryl ether sulphate (2 EO) (27—28% WAS)	60	60	60	60	60
Sodium lauryl sulphate (over 90% WAS)	ž	5	5	5	5
Isopropyl myristate	****	5			••••
(A)		-	5	••••	
(B)		****		5	••••
(C)	****	p.0, 0 0.0	****	****	5
Water	35	30	30	30	30
Appearance	clear	turbid deposit formed	clear	clear	clear
Foaming power					
Initial volume in m	il 1 minute a	fter beating			
0.5 g/litre	260	170	300	340	240
1.0 g/litre	510	250	490	490	400
2.0 g/litre	660	330	610	640	540
Breaking down of t	the volume of	foam in ml/min	ute		
0.5 g/litte	2.0	2.5	4.0	4.5	1.5
1.0 g/litre	5.5	3.5	7.0	8.8	4.0
2.0 g/litre	8.0	6.5	9.5	3.0	6.5

The foaming power of the individual mixtures was measured in the foam-beating machine according to DIN (Deutsche Industrie Norm) classification No. 53,902, in which the volume of foam was measured. The figures for the foam were taken at 45°C in water of 10° German hardness after 30 beats. The measurement was taken I minute after the end of the beating and 21 minutes after the end of the beating. The breakdown of the volume of foam was calculated in ml/minute from the decrease of the volume of foam in

20 minutes. The amounts given in g/litre relate to the respective mixture.

As may be seen from the above Table, the foam values of the mixtures containing oil replacement means according to the invention are substantially better than when isopropyl myristate is used as oil replacement materials, and are scarcely inferior to the figures for a mixture of pure detergent substances.

A few formulations for cosmetic preparations containing oil replacement materials according to the invention are given below.

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Clear Shampoo			******	
Sodium lauryl sulphate (2EO)				
(2728% WAS)	40	parts	hv	weigh
Coconut fatty acid diethanolamide	6	72	22	55
Oil replacement means (A)	10	52	33	35
Water	44	92	23	22
Shampoo for dry hair				
Sodium lauryl ether sulphate				
(2 EO) (27—28% WAS) Sodium lauryl sulphate	20	parts	by 1	weight
(90% WAS) Coconst fatty acid diethanolamide	5	23	22	35
Coconut fatty acid monoethanol-	3	32	33	2.7
amide paste 30%	5	35	59	2.r 22
Water-soluble vitamin F	0.5	25	52	35
Oil replacement means (B)	25.0	35	3>	25
Water	41.5	35	23	33
Foam bath				
Sodiem lauryl ether sulphase				
(2 EO) (27—28% WAS) Sodium lauryl sulphase	30 pa	rts by	wci	ght
(90% WAS)	15 ,	£ .	23	
Coconut facty acid diethanolamide	ž.,	2 25	35	
Pine-needle oil		55	53	
Oil replacement means (C) Water	10 , 35 ,	205 20 20	20 22	
Hair wash				
Isopropanot	60.0	parts l	***	rains
Menthol	0.2			
Calcium pantothenate	0.05		53	22
Vitamin H	0.30		>> >>	23
Inosital	0.10		KS	52
Perfume	0.50	39 :	kt	22
Oil replacement means (C)	5.00	39	22	25
Water	33.85	22 :	**	539
After-shave lotion				
Ethyl alcohol 96%	65 0	parts b	y w	eight
Menthol	0.2		"	22
Camphor Peruvian balsam	0.2	3 2 3	25	35
retuvian baisam Perfume	0.1	, פל.	25	33
Citycerine	9.5 5.8	33 3	12	3.2
Witch hazel extract	10.0		Ú5	22
Boric scid	0.5		73	>>
Oil replacement means (A)	10.0		15 15	35
NTT		40 3		-17

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Sun tan cream

Colloidally dispersed mixture of 90 parts of cetylstearyl alcohol and 10 parts of sodium lauryl sulphate 2-Octyldodecanol Groundnut oil Light-protective means Oil replacement means (B) Water

10.0 parts by weight 10 5 2 35 20 ,9,9 22 53

The oil replacement compositions according to the invention can be used particularly advantageously in cosmetic cleansing means, because they do not exert any appreciable influence on the foaming power of the surface-active products, and because they already have a good solubility in alcoholwater mixtures.

WHAT WE CLAIM IS:-

1. A composition, for use in cosmetic preparations, which comprises a surface active compound and the esserification product of an ethylene oxide addition compound produced from glycerine and 4 to 20 mol of ethylene oxide per mol of glycerine with a faity acid of chain length from 8 to 18 carbon atoms in a ratio of 1 to 2 mol of fatty acid to 1 mol of glycerine-ethylene oxide addition compound as oil replacement material.

2. A composition according to claim 1, in which the ratio of glycerine to ethylene oxide in the addition compound is from 7 to 15 mols of ethylene oxide per mol of glycerine.

3. A composition according to claim 1 or 2 wherein the fatty acids of chain length from 8 to 18 carbon atoms are in a ratio of 1 mel of fatty acid to 1 mol of the glycerine-ethylene exide addition compound.

4 A composition according to claims 1 to 3, containing from 2 to 50% by weight of

the oil replacement material.

5. A composition according to claim 1 to 4 comaining from 5 to 25% by weight of the oil replacement material.

6. A composition according to claim 1 substantially as hereinbefore described with reference to and as illustrated in the foregoing examples.

7. A cosmetic preparation whenever containing a composition as claimed in any one

of claims 1 to 6. W. P. THOMPSON & CO., 12, Church Street, Liverpool, LI BAB. Chartered Patent Agents.

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